

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
28 December 2000 (28.12.2000)

PCT

(10) International Publication Number
WO 00/78286 A1

(51) International Patent Classification⁷: **A61K 9/12, B65D 83/14**

(21) International Application Number: **PCT/US00/16462**

(22) International Filing Date: **15 June 2000 (15.06.2000)**

(25) Filing Language: **English**

(26) Publication Language: **English**

(30) Priority Data:
60/139,961 **18 June 1999 (18.06.1999) US**

(71) Applicant: **3M INNOVATIVE PROPERTIES COMPANY [US/US]; 3M Center, P.O. Box 33427, Saint Paul, MN 55133-3427 (US).**

(72) Inventors: **WU, Zheng, Z.; P. O. Box 33427, Saint Paul, MN 55133-3427 (US). GOVIND, Nayna; P.O. Box 33427,**

Saint Paul, MN 55133-3427 (US). JOHNSTON, Peter, R.; P.O. Box 33427, Saint Paul, MN 55133-3427 (US).

(74) Agents: **RINGSRED, Ted, K. et al.; Office of Intellectual Property Counsel, P.O. Box 33427, Saint Paul, MN 55133-3427 (US).**

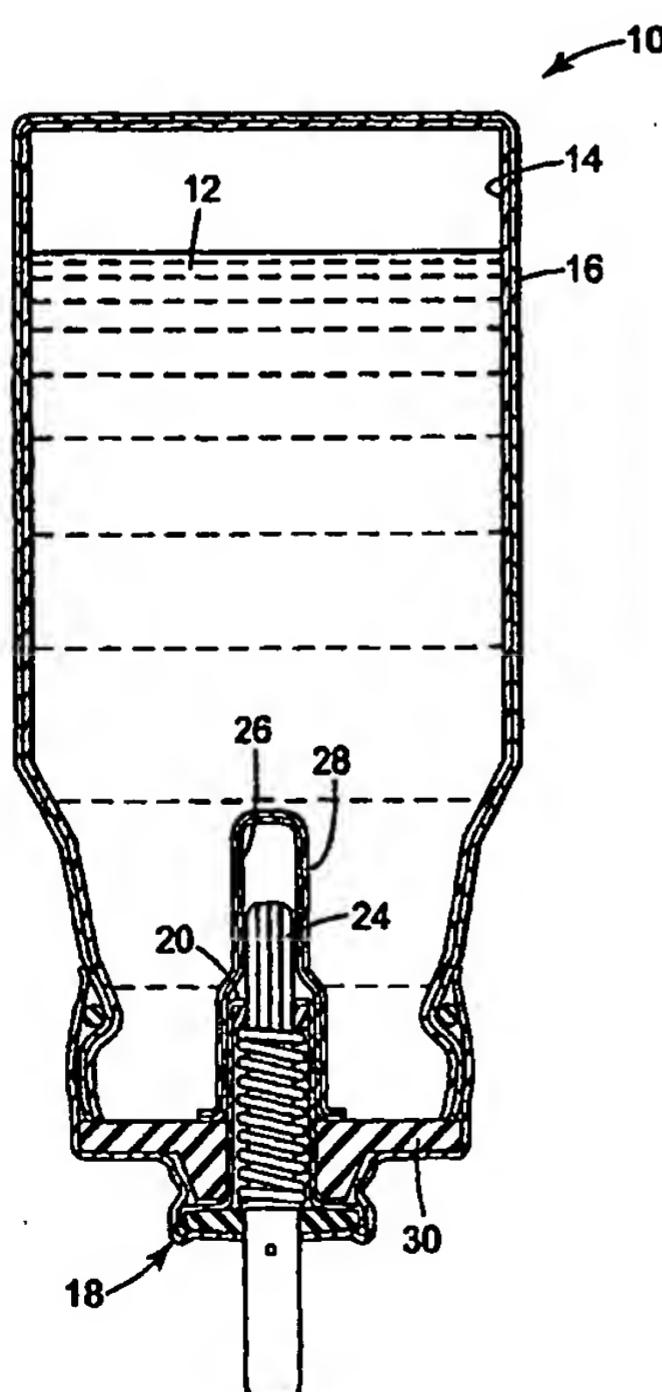
(81) Designated States (*national*): **AE, AG, AL, AM, AT, AT (utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, CZ (utility model), DE, DE (utility model), DK, DK (utility model), DM, DZ, EE, EE (utility model), ES, FI, FI (utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KR (utility model), KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (utility model), SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.**

(84) Designated States (*regional*): **ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European**

[Continued on next page]

(54) Title: **STEROID SOLUTION AEROSOL PRODUCTS WITH ENHANCED CHEMICAL STABILITY**

(57) Abstract: A medicinal aerosol steroid solution formulation product with enhanced chemical stability. The steroid is a 20-ketosteroid having an OH group at the C-17 or C-21 position and the aerosol container has a non-metal interior surface which has been found to reduce chemical degradation of such steroids.



WO 00/78286 A1



patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

— Before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments.

Published:

— With international search report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

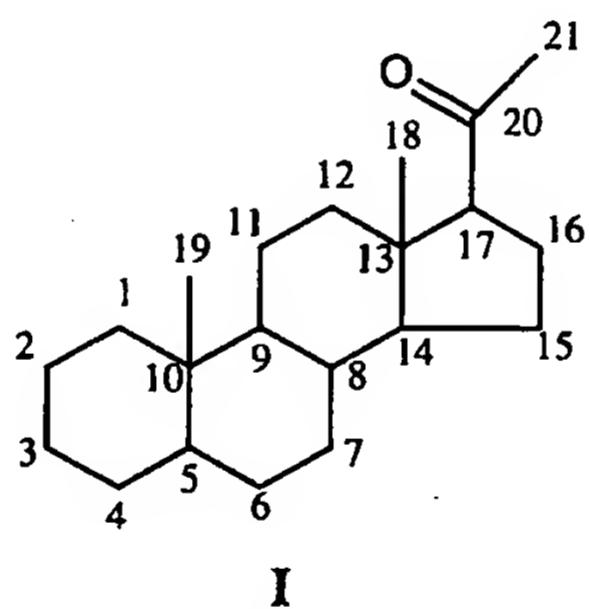
STEROID SOLUTION AEROSOL PRODUCTS
WITH ENHANCED CHEMICAL STABILITY

5 This application claims the benefit of earlier filed U.S. provisional application 60/139,961, filed on June 18, 1999.

The present invention relates to medicinal aerosol products and, in particular, to aerosol products such as metered dose inhalers for delivery of steroids. The invention is particularly related to certain 20-ketosteroids that have been found to be highly susceptible 10 to chemical degradation when formulated as solution aerosol products, and provides a way of enhancing chemical stability of such steroids.

Structure I shown below is a typical core structure for a large number of natural and synthetic 20-ketosteroids, with the standard IUPAC numbering system of the carbon positions 1 to 21 indicated.

15



These types of steroids, with varying substituents and bonding, have many well-known therapeutic uses, especially based upon their anti-inflammatory activity. It is often desirable to deliver such steroids topically using aerosol spray devices, such as metered 20 dose inhalers (MDIs). MDIs are commonly used to deliver steroids, e.g., beclomethasone dipropionate, to the airways of patients via oral or nasal inhalation for the treatment of asthma and allergic rhinitis.

One common difficulty, however, in making products for delivering steroids has been that they are often chemically unstable in aerosol formulations and degrade during 25 storage. A great deal of research has been directed at steroid degradation. Chemical degradation is especially problematic when the steroid is dissolved in the formulation and,

consequently, the vast majority of marketed MDI steroid products are formulated as particulate suspensions of the steroid, which are much less susceptible to chemical degradation than solutions. For example, it is believed that all currently marketed CFC-containing MDI products for delivering steroids are available only as particulate suspension formulations in CFC propellants.

More recently, some selected steroids have been reformulated as solutions in non-CFC hydrofluorocarbon (HFC) propellants with ethanol. In the case of beclomethasone dipropionate, for example, solution formulations are disclosed in U.S. 5,766,573 which are surprisingly chemically stable in propellant HFC 134a and/or 227 and ethanol in a conventional aluminum canister. Likewise, ciclesonide formulations are surprisingly chemically stable in certain solution MDI formulations disclosed in WO 98/52542. Solution formulations of flunisolide are disclosed in U.S. 5,776,433, where it is indicated that chemical stability may be enhanced by using additives like water, sorbitan trioleate, and cetylpyridinium chloride, and also that certain containers such as glass and resin coated aluminum enhance chemical stability and/or minimize the absorption of flunisolide onto the container wall. Also, WO 96/40042 discloses that aqueous formulations of triamcinolone acetonide in neutral or basic solutions undergo oxidative degradation catalyzed by trace levels of metal ions, especially copper, and proposes the use of EDTA as sequestering agent and/or adjusting pH.

Despite these limited examples, though, most commercial MDI formulations of steroids and other drugs have continued to be particulate suspensions. WO 98/13031, for example, discloses recent work on reformulating budesonide as a non-CFC suspension formulation. However, suspension formulations of a medicament are more likely to encounter problems with physical instability (e.g., agglomeration, crystal growth and deposition on the container wall, all resulting in inconsistent dosage delivery).

A drug delivery device providing medicinal steroid solution formulations with enhanced chemical stability would offer some significant advantages over suspension formulations. Besides homogeneity, solution formulations have been found in some cases—e.g., using HFC propellants and low ethanol content--to give dramatically higher respirable fractions (i.e., the percentage of active ingredient able to reach the airways of the lung) compared to a particulate suspension of the steroid drug. See U.S. 5,776,432. Furthermore, an aerosol product providing a chemically and physically stable aerosol

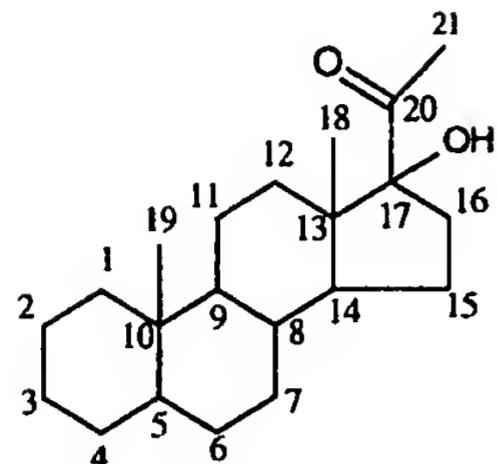
steroid formulation using non-CFC propellant would offer the advantage of being more ozone friendly than currently available aerosol products with CFCs.

Nevertheless, despite a substantial need, the problem of chemical degradation in steroid solution aerosol products has been poorly understood. Until now there has been no
5 way to identify which steroids are likely to be most stable as solution aerosols and which will be most sensitive to degradation in solution aerosol products or how to reduce such degradation.

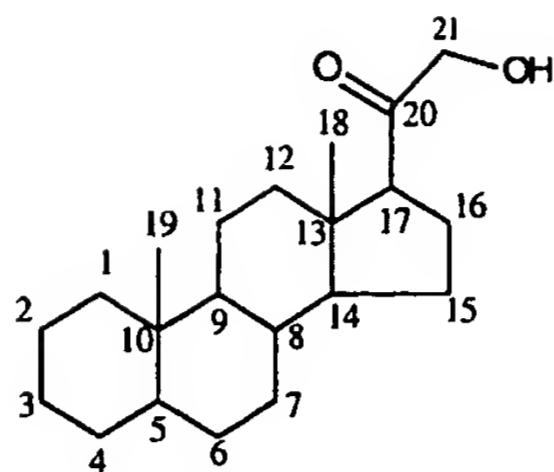
It has now been found that those steroids in particular having a C-20 ketone and an OH group at the C-17 position or, especially, the C-21 position or both (hereafter
10 collectively referred to as "C-17/21 OH 20-ketosteroids") are subject to enhanced chemical degradation when stored in contact with a metal container (particularly the metal oxide e.g., Al₂O₃ layer that forms on the interior surface of the container). Moreover, the vast majority of MDI's on the market, including all MDI steroid products, use aerosol
15 containers made of metal, usually aluminum. By utilizing an aerosol container having a non-metal interior surface it is possible to produce solution aerosol formulations of C-17/21 OH 20-ketosteroids having enhanced chemical stability.

According to the invention there is provided a medicinal aerosol steroid solution formulation product with enhanced chemical stability, comprising an aerosol container equipped with a dispensing valve and containing a medicinal aerosol formulation having a
20 20-ketosteroid drug dissolved therein, said 20-ketosteroid having an OH group at the C-17 or C-21 position or both, with the proviso that said 20-ketosteroid is other than flunisolide, and wherein said container is provided with a non-metal interior surface so as to reduce chemical degradation of the 20-ketosteroid.

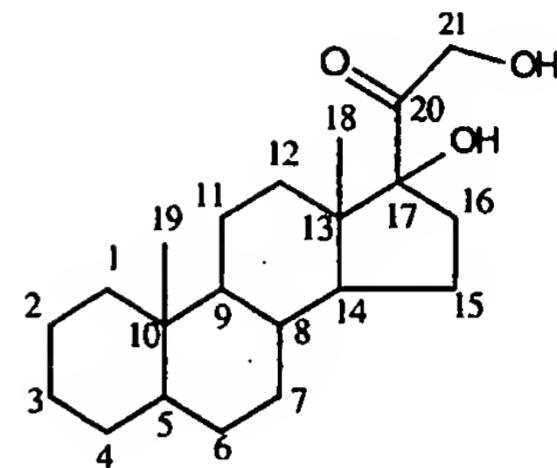
Generic structures for typical C-17 OH, C-21 OH, and C-17 and 21 OH 20-
25 ketosteroids are shown below in structures II-IV, respectively.



II



III



IV

The present invention is especially preferred with respect to C-21 OH 20-ketosteroids (with or without a C-17 OH group). The C-21 OH group can substantially increase biological activity of a steroid, but such steroids are also much more susceptible to chemical degradation in the presence of metal. Particularly preferred 20-ketosteroids are budesonide, triamcinolone acetonide, dexamethasone, and betamethasone 17-valerate, all of which have an OH group at the C-21 position. These steroids are all currently on the market as CFC particulate suspension formulations in metered dose inhalers and, at least in the case of budesonide, work has been conducted to reformulate this important steroid as a non-CFC suspension product in HFA propellants. See PCT published application WO98/13031.

The most preferred type of container for use in the present invention is a conventional aluminum (or aluminum alloy) aerosol canister, but with an interior coating of an inert material, such as a spray-coated, baked epoxy-phenolic lacquer (available from Cebal Printal U.K. Ltd.). Other metals, such as stainless steel, may likewise be used with an inert interior coating. It is also preferred that the internal surfaces of metal valve components in contact with the formulation are similarly coated with an inert material. Another preferred coating for the inside of the container is perfluoroethylenepropylene (FEP).

A preferred coating for the metal valve components is a very thin layer of glass, or other material, deposited by gas vapor deposition. Such coating is preferably used on all of the metal valve components in contact with the formulation, including the inside and outside of the metering chamber, inside and outside of the bottle emptier (if any), and the inside and outside of the valve stem (if metal), and may also be used to coat the inside of

the canister. The preferred such coating technique is the Silcosteel™ process available from Restek Corporation, Bellefonte, PA. The Silcosteel™ aspect of the invention is useful even outside the context of the chemical degradation problem, for both solution and suspension formulations.

5 Preferred formulations use a liquified propellant such as hydrogen-containing (non-CFC) propellants, more preferably hydrofluorocarbons, such as 134a and/or 227. Particularly preferred formulations include about 0.1 to 0.5% C-17/21 OH 20-ketosteroid about 75 to 99% 134a and/or 227, and about 1 to 25 % w/w ethanol, more preferably about 80 to 95% 134a and/or 227, and about 5 to 20% ethanol. The most preferred medicinal
10 aerosol products according to the invention are MDI's comprising about 0.1 to 0.5% budesonide or triamcinolone acetonide dissolved in about 80-95% 134a and/or 227 and about 5-20% ethanol, contained in a coated aluminum aerosol canister equipped with a metering valve.

15 It should also be noted that chemical stability is especially critical for MDIs since these medicinal aerosol products must remain stable and deliver accurate dosing throughout their shelf life (typically 2 to 3 years) and in use. Only a very small amount of chemical degradation can be tolerated. Moreover, by providing chemically stable solutions of C-17/21 OH 20-ketosteroids, such as budesonide, triamcinolone acetonide, dexamethasone, and betamethasone 17-valerate, MDIs can be made that produce extra-
20 fine aerosols resulting in higher respirable fractions than suspension formulation products.

It can thus be seen that the present invention provides a medicinal aerosol steroid solution formulation product with enhanced chemical stability. Such product includes a container equipped with a dispensing valve and containing a medicinal aerosol formulation having a 20-ketosteroid drug dissolved therein. The 20-ketosteroid is other
25 than flunisolide and has an OH group at the C-17 or C-21 position or both, and the container is provided with a non-metal interior surface so as to reduce chemical degradation.

Also provided is a method of reducing the chemical degradation of a medicinal 20-ketosteroid dissolved in a formulation contained in a metal container, said 20-ketosteroid
30 being other than flunisolide and having an OH group at the C-17 position or C-21 position or both, comprising the step of providing a coating of inert material on the interior surface

of the metal container so as to reduce reaction of the 20-ketosteroid with metal oxides from the container.

The invention further provides a process for making a chemically stable 20-ketosteroid solution aerosol product, by filling into a container an aerosol formulation comprising a dissolved 20-ketosteroid other than flunisolide, said 20-ketosteroid having an OH group at the C-17 position or C-21 position or both, and said container having an inert non-metal interior surface so as to avoid chemical degradation of the 20-ketosteroid due to interaction with the container.

The invention will be described with reference to the following drawing, wherein:
10 Fig. 1 is a cross-sectional view of a metered dose inhaler containing a medicinal 20-ketosteroid formulation with enhanced chemical stability according to the present invention;

Fig. 2 is the same as Fig. 1, but with a modified valve configuration.

Turning to Fig. 1, there is shown a medicinal aerosol device 10 comprising a pressurizable metal aerosol container 16 equipped with a metering valve 18. The metal container 16 is preferably made of aluminum (i.e. aluminum or aluminum alloy) and has an inert interior coating layer 14. The metering valve 18 includes a metal metering chamber 20 with a coating layer 22. Moreover, although not shown, it is preferred that as many other metal surfaces in contact with the formulation 12 as feasible are also coated with an inert layer (e.g., the interior of the metering chamber). It is also preferred to use a valve design that minimizes metal surfaces in contact with the formulation 12. For example, it is preferred in the context of the present invention to use a plastic valve stem 24 instead of metal.

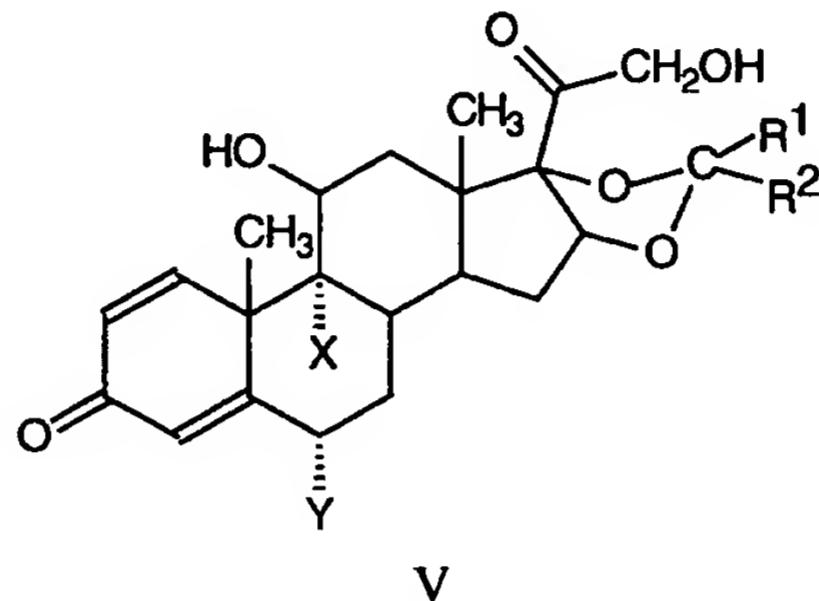
Fig. 2 shows an alternative preferred embodiment that is essentially the same as Fig. 1, but utilizes a fixed bottle emptier 26, with coating layer 28. Also, a solution gasket 30 is used to further prevent contact of the formulation with metal components.

The medicinal aerosol formulation 12 preferably includes a non-CFC propellant, a cosolvent (if necessary), and a therapeutically effective amount of dissolved C-17/21 OH 20-ketosteroid. A therapeutically effective amount will normally be a concentration so as 30 to provide in the range of about 100 to 1500 mg per day using one to eight puffs.

Preferred propellants include hydrogen containing propellants, such as HFCs 134a and/or 227. Ethanol is a preferred cosolvent, although other cosolvents and solubilization aids

(e.g., surfactants) may be used. The amount of cosolvent used is preferably an amount sufficient to completely dissolve the 20-ketosteroid. The formulation 12 may also include other excipients, such as stabilizers, antioxidants, flavoring agents, and the like.

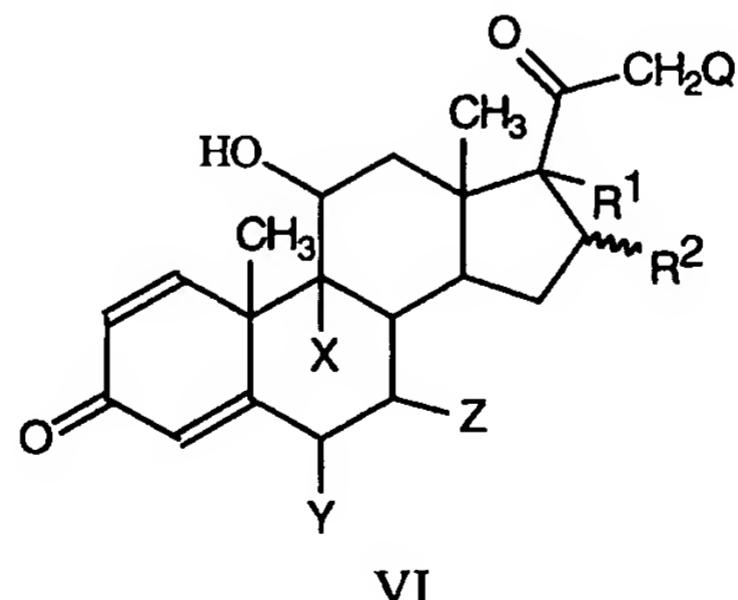
5 Although there are many possible C-17/21 OH 20-ketosteroids, there are two preferred types set forth below as structures V and VI:



Wherein X is H, Cl, or F; Y is H, F, or Me; R¹ is H or an alkyl group; R² is an alkyl group; and provided that when Y is F, then X is not H. Preferred R¹ and/or R² alkyl groups are methyl, ethyl, propyl, butyl, pentyl, and hexyl groups, including their branched and cyclic isomers. R¹ may form a ring with R², preferably a cyclopentyl or cyclohexyl ring.

10 Examples of known 20-ketosteroids according to structure V include budesonide, triamcinolone acetonide, desonide, and fluocinolone acetonide.

15



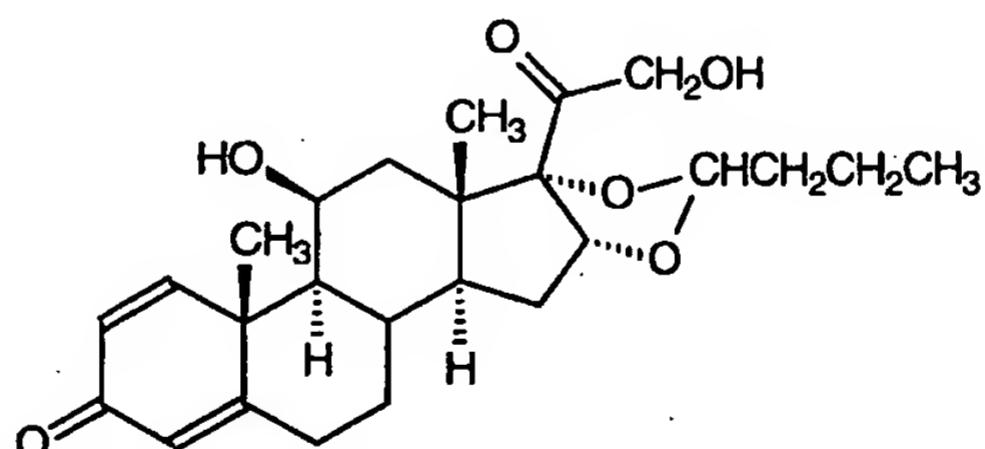
Wherein Q is OH, H, Cl, or PO(ONa)₂; X is H, Cl, or F; Y is H, F, or Me; Z is H or Cl; R¹ is H, OH or propionate provided that when Q is not OH, then R¹ must be OH; and R² is H, OH, or Me.

Examples of known 20-ketosteroids according to structure VI include alclometasone, beclomethasone, beclomethasone 17-monopropionate, betamethasone, betamethasone 17-valerate, clocortolone, desoximetasone, dexamethasone, dexamethasone sodium phosphate, dexamethasone 21-isonicotinate, diflorasone, flumethasone, 5 methylprednisolone, paramethasone, prednisolone, tramcinolone, clobetasol, and fluorometholone.

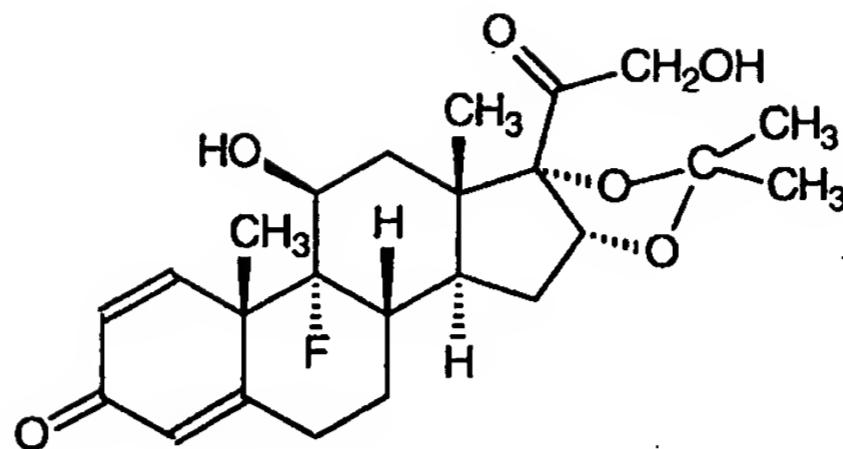
Within the group of C-17/21 OH 20-ketosteroids, those having an OH group at C-21, with or without an OH group at C-17, suffer from more severe degradation in the presence of metal oxide than 20-ketosteroids having an OH group at C-17, but not at C-21. 10 For example, beclomethasone 17-monopropionate (17-BMP, 21-OH) degrades in ethanol/Al₂O₃ about 100 times faster than its isomer 21-BMP (17-OH). This is important since it has been reported that the Structure-Activity Relationship for a C-21 OH group has an enhancement factor toward anti-inflammatory activity of 25 compared to an enhancement factor of only 4 for a C-17 OH group. See Doerge, R.F., Wilson and 15 Grisvold's Textbook of Organic Medicinal and Pharmaceutical Chemistry, 8th Edition, p. 703, table 18-13 (J.B. Lippincott Co., Philadelphia, 1982).

For particular structural examples, four preferred C-17/21 OH 20-ketosteroids according to the present invention, budesonide, triamcinolone acetonide, betamethasone 17-valerate, and dexamethasone, have the following structures:

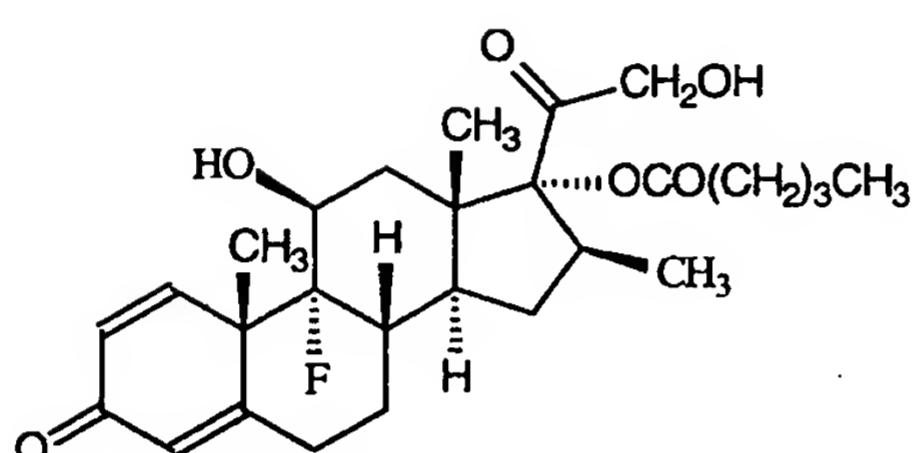
20



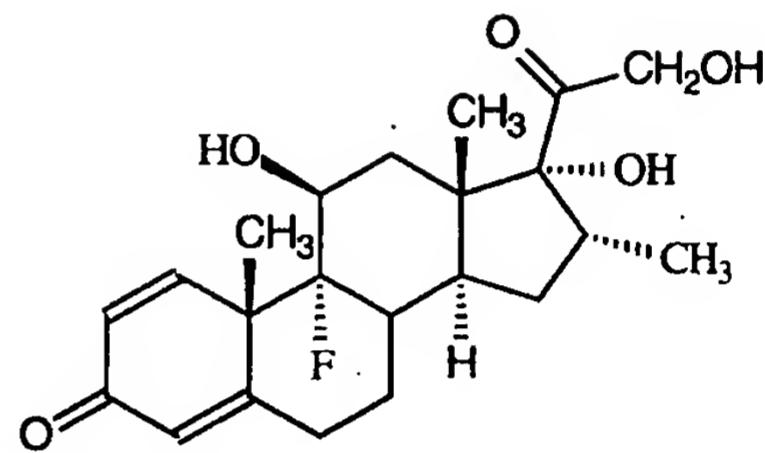
Budesonide



Triamcinolone Acetonide



Betamethasone 17-Valerate



Dexamethasone

5 The C-17/21 OH 20-ketosteroid is dissolved in the aerosol formulation and
preferably contained in an inertly coated metal container. As used herein, the terms
“coated”, “inert coating” and the like simply refer to a non-metal interior coating that does
not promote degradation at the OH substituents on C-17/21 OH 20-ketosteroids. Inert
coating materials include any suitable polymer, lacquer, resin, or other coating treatment
10 that creates a barrier to chemical interaction of the dissolved C-17/21 OH 20-ketosteroid
and metal on the container (especially metal oxides).

15 Examples of suitable interior coatings include epoxy-phenolic resins, epoxy-urea-
formaldehyde resins, polytetrafluoroethylene (PTFE), perfluoroethylenepropylene (FEP),
perfluoroalkoxyalkane (PFA), ethylene tetrafluoroethylene (ETFE), poly(vinylidene
fluoride) (PVDF), and chlorinated ethylene tetrafluoroethylene. Blends may be used of
fluorinated polymers with non-fluorinated polymers such as polyamides, polyimides,
polyethersulfones, polyphenylene sulfides, and amine-formaldehyde thermosetting resins.
Specific blends include PTFE/FEP/polyamideimide, PTFE/polyether sulphone (PES) and
FEP-benzoguanamine. Preferred interior coating materials are epoxy-phenolic resins,
20 epoxy-urea-formaldehyde resins, PTFE, and FEP. Additional information regarding

interior can coatings is taught in, e.g., EP 642992, WO 96/32099, WO 96/32150, WO 96/32151, and WO 96/32345, which disclose interior can coatings for drug suspension formulation products.

A preferred coating for the metal valve components is a very thin layer of fused silica glass, or other material, deposited by gas vapor deposition. Such coating is preferably used on all of the metal valve components in contact with the formulation, including the inside and outside of the metering chamber, inside and outside of the bottle emptier (if any), and the inside and outside of the valve stem (if metal). The preferred such coating technique is the Silcosteel™ process available from Restek Corporation, Bellefonte, PA. This process deposits a submicron layer of fused silica glass on the metal components and can be used both on the valve components and on the interior of the canister. Not only is it helpful in preventing chemical reaction with the metal, but passivation of the metal surface using the Silcosteel process can provide a smooth surface on, for example, the valve stem so as to reduce friction and help prevent valve clogging, and can also reduce oxidation of the metal that can introduce particulate material into the system. The Silcosteel™ aspect of the invention is thus useful even outside the context of the chemical degradation problem, for both solution and suspension formulations. The Silcosteel process is performed at a temperature of 400°C, which has the added benefit of thermally removing residual oils on the metal surface.

20

It is also possible to use containers made from non-metal materials, such as glass or plastic (e.g., polyethylene terephthalate, from Precise Plastic Ltd., U.K.). In the case of glass containers, however, it is preferred to use glass having a low metal oxide content. According to the American Society for Testing and Materials (ASTM), Type I/Class A glass contains 2% aluminum oxide, whereas Type II/Class B glass contains 7% aluminum oxide. The former is therefore preferred. Type III (soda-lime) glass vials may also be used (available from Wheaton Coated Products). High metal oxide content glass should also be avoided during development and testing of formulations containing C-17 and/or C-21 OH 20-ketosteroids because such use could cause unsuspected chemical degradation due to the glass metal oxide content.

Coated containers can be made by pre-coating the metal roll stock before forming the container or by coating the container after it is made, by various techniques known in

the art of coating. For example, suitable coating procedures include plasma coating, electrostatic dry powder coating, impregnating/spraying, hard anodization with polymer deposition, chemical vapor deposition (CVD), physical vapor deposition (PVD), as well as other procedures known in the art for this purpose. Preferably, the containers are coated
5 with an epoxy resin and then baked. Suitable coated containers can be obtained from, for example, Cebal Printal U.K. Ltd.

Preferred products/devices according to the present invention are pressurized aerosols such as MDIs that use liquefied gas propellants, including chlorofluorocarbons (CFCs), hydrofluorocarbons (HFCs), fluorocarbons (FCs), hydrocarbons (HCs),
10 hydrochlorofluorocarbons (HCFCs), and dimethyl ether (DME). Propellants containing hydrogen are preferred. Ethanol may also be included to assist in solubilizing the 20-ketosteroid, preferably in an amount of about 1-25%. Formulations comprising dissolved C-17/21 OH 20-ketosteroid, HFC propellant, such as 134a and/or 227, with 5-20% ethanol are particularly preferred. Also, it is generally preferred that the formulations are
15 essentially non-aqueous, meaning that they do not include added water, although very small amounts of water may be present due to water ingress and/or as a residual in formulation components (such as ethanol). Formulations will preferably have less than about 3%, and more preferably less than about 0.5 % water content.

The containers can be equipped with any suitable conventional or unconventional
20 dispensing valve, preferably a metered dose valve.

The following examples illustrate various aspects of the invention.

Example 1

The following illustrates the effect of a metal oxide (Al_2O_3) in the degradation of a
25 C-17/21 OH 20-ketosteroid, in this case budesonide, which has an OH group attached on the C-21 carbon position.

At ambient temperature, a 3 mL ethanol solution of budesonide (0.007 M) was mixed with 0.1 g of aluminum oxide (Aldrich, neutral, activated Brockmann I, 150 mesh), which mimics aluminum oxide existing on the inner surface of aluminum containers. The
30 mixture was held at 75°C for several days, and an aliquot was taken at each of the time intervals given in the Table 1. Each aliquot was filtered with a syringe filter and diluted with acetonitrile by 20 times for HPLC analysis to determine the recovery of budesonide

with respect to the initial concentration. The recoveries of budesonide were compared with a reference, which contained the same concentration of budesonide but in ethanol solution without aluminum oxide. The results are summarized in Table 1.

5 Table 1. Recovery (%) of budesonide in a mixture of ethanol/aluminum oxide at 75°C or in ethanol at 75°C.

Time (Hours)	In Ethanol/Aluminum Oxide	In Ethanol
0	100	100
6	4.97	-
24	0.20	99.2
48	0.22	99.4
168	-	95.5

10 In the presence of aluminum oxide, budesonide was almost totally consumed in 6 hours at 75°C. In contrast, without aluminum oxide the recovery of budesonide was as high as 95.5% in 168 hours at the same temperature. The data demonstrate that the presence of aluminum oxide substantially facilitates the decomposition of budesonide. Ethanol, however, appears to play only a minor role in the decomposition.

Example 2

15 The following Example illustrates the effect of aluminum oxide in the degradation of another C-17/21 OH 20-ketosteroid, in this case triamcinolone acetonide, which has an OH attached to the C-21 carbon position.

20 At ambient temperature, a 3 mL ethanol solution of triamcinolone acetonide (0.007 M) was mixed with 0.1 g of aluminum oxide (Aldrich, neutral, activated Brockmann I, 150 mesh), which mimics aluminum oxide existing on the inner surface of aluminum containers. The mixture was held at 75°C for several days, and an aliquot was taken at each of the time intervals given in the Table 2. Each aliquot was filtered with a syringe filter and diluted with acetonitrile by 20 times for HPLC analysis to determine the recovery of triamcinolone acetonide with respect to the initial concentration. The 25 recoveries of triamcinolone acetonide were compared with a reference, which contained

the same concentration of triamcinolone acetonide but in ethanol solution without aluminum oxide. The results are summarized in Table 2.

5 Table 2. Recovery (%) of triamcinolone acetonide in a mixture of ethanol/aluminum oxide at 75°C or in ethanol at 75°C.

Time (Hours)	In Ethanol/Aluminum Oxide	In Ethanol
0	100	100
6	70.2	-
24	19.6	98.6
48	12.4	98.5
168	-	97.5

10 In the presence of aluminum oxide, triamcinolone acetonide recovery was 12.4% in 48 hours at 75°C. In contrast, without aluminum oxide the recovery of triamcinolone acetonide was as high as 97.5% in 168 hours at the same temperature. The data demonstrate that the presence of aluminum oxide substantially facilitates the decomposition of triamcinolone acetonide. Ethanol, however, appears to play only a minor role in the decomposition.

15

Example 3

The following Example illustrates the effect of aluminum oxide in the degradation of C-17/21 OH 20-ketosteroid, in the case dexamethasone, which has one OH attached to C-21 and another OH attached to C-17.

20

At ambient temperature, a 3 mL ethanol solution of dexamethasone (0.007 M) was mixed with 0.1 g of aluminum oxide (Aldrich, neutral, activated Brockmann I, 150 mesh), which mimics aluminum oxide existing on the inner surface of aluminum containers. The mixture was held at 75°C for several hours, and an aliquot was taken at each of the time intervals given in Table 3. Each aliquote was filtered with a syringe filter and diluted with acetonitrile by 20 times for HPLC analysis to determine the recoveries of dexamethasone with respect to the initial concentration. The recoveries of dexamethasone were compared

25

with a reference, which contained the same concentration of dexamethasone but in ethanol solution without aluminum oxide. The results are summarized in following Table 3.

5 Table 3. Recovery (%) of dexamethasone in a mixture of ethanol/aluminum oxide at 75°C or in ethanol at 75°C.

Time (hours)	in Ethanol/Aluminum Oxide	in Ethanol
0	100	100
3	6.0	-
6	4.0	-
10	0.0	-
24	-	98.0
72	-	96.0
168	-	90.0

10 In the presence of aluminum oxide, dexamethasone completely degrades after ten hours at 75°C. In contrast, without aluminum oxide the recovery of dexamethasone is as high as 90% after 168 hours at the same temperature. The data demonstrate that aluminum oxide substantially facilitates degradation of dexamethasone in ethanol solution. Ethanol, however, appears to play a only a minor role in degradation.

Example 4

15 Table 4 below shows results of solubility testing in mg/g of budesonide in P134a and P227 formulations at approximately 5 °C.

Table 4. Solubility of budesonide

Sample	Budesonide Solubility mg/g (5°C)		
	1	2	MEAN
P134a	0.013	0.010	0.012
P134a/5% EtOH	0.459	0.471	0.465
P134a/10% EtOH	1.370	1.394	1.382
P134a/ 15% EtOH	2.686	2.711	2.699
P134a/20% EtOH	4.329	4.390	4.360
P227	0.017	0.015	0.016
P227/5% EtOH	0.652	0.661	0.657
P227/10% EtOH	1.698	1.707	1.703
P227/15% EtOH	2.952	3.008	2.98
P227/20% EtOH	4.387	4.483	4.435

Examples 5-6

5 Experiments were conducted to test the influence of various factors on chemical stability of two exemplary budesonide solution formulations, examples 5 and 6, shown in Table 5.

Table 5. Exemplary budesonide formulations

10

Example 5	% w/w
Budesonide	0.362
Oleic acid	0.029
Ethanol absolute	20.000
Propellant 134a	79.609
Example 6	%w/w
Budesonide	0.327
Oleic acid	0.029
Ethanol absolute	20.000
Propellant 227	79.644

15

The above formulations of Examples 5 and 6 were manufactured on a pilot-scale cold filler and filled into metered dose inhaler vials having various different components: different valve types, coated versus uncoated aluminum cans, and with and without a nylon desiccant insert. The units were then inverted and stored at either 40°C/85% RH for 14 days or 30°C/30% RH for 17 days prior to analysis of budesonide degradation.

The test results indicate that epoxy-phenolic lacquer-coated aluminum containers (from Cebal Printal U.K. Ltd.) showed in all but one case significantly enhanced chemical stability versus formulations in uncoated cans.

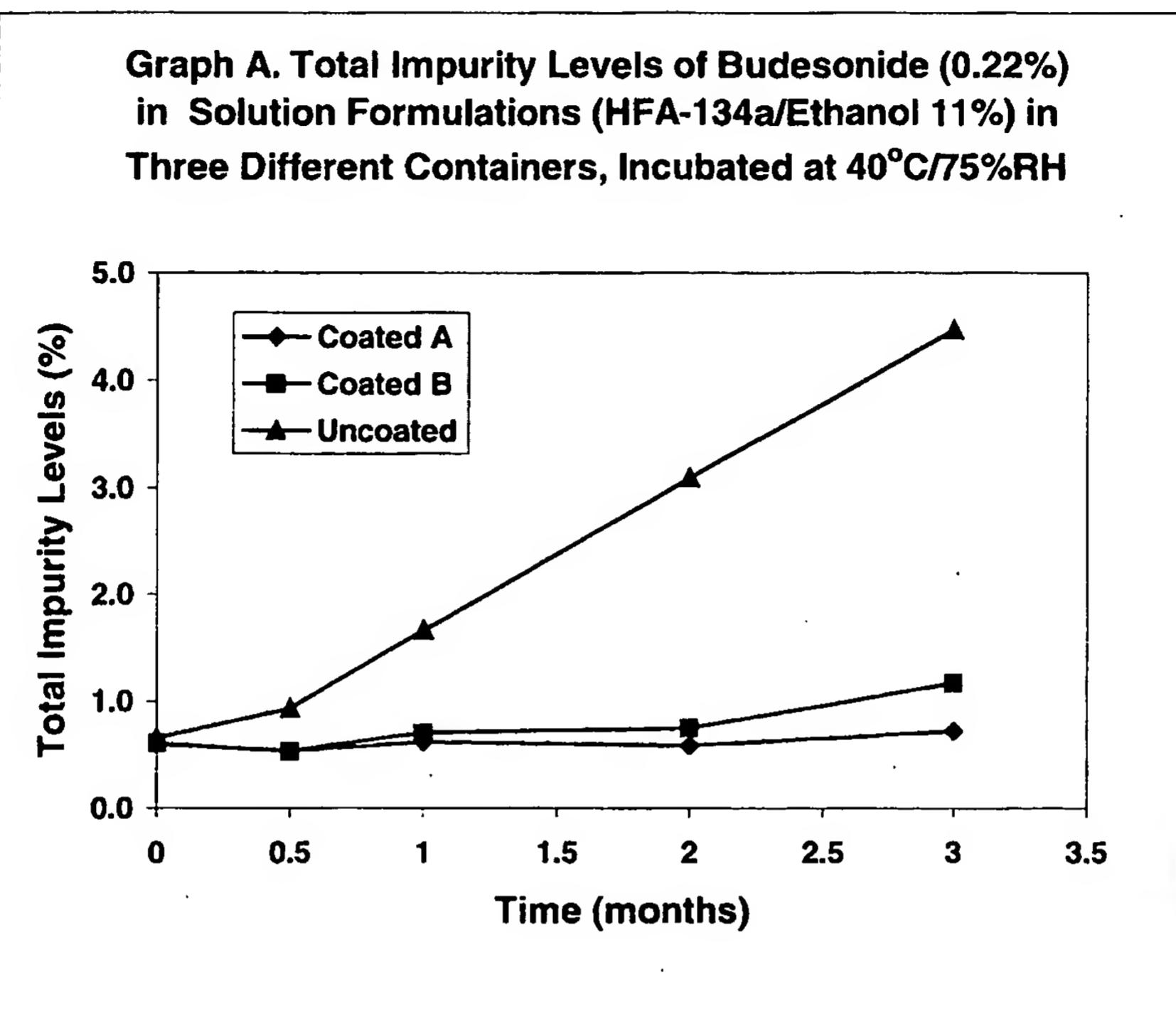
5 The type of valve used also had an impact on chemical stability. The greatest chemical stability enhancement was found in units using a Bespak 357 valve with EPDM seals, a Cebal epoxy lacquered can, and the formulation of Example 6. Nearly as stable were units equipped with a 3M Neotechnic Spraymiser™ HFA valve, a Cebal epoxy lacquered can, and the formulations of Examples 5 or 6. Also, none of the valves tested were coated and it is expected that this would further enhance chemical stability.

10 Other factors also appeared to affect chemical stability. Formulations using propellant 227 appeared to be slightly more stable. Containers having a desiccant generally exhibited reduced degradation compared to those without desiccant inserts. This suggests that water may increase degradation and that using dried ethanol may thus help further enhance chemical stability. Also, the use of certain antioxidants such as ascorbic acid and ascorbyl palmitate (but not vitamin E) appeared to enhance chemical stability,
15 while the use of oleic acid appeared to reduce chemical stability.

Example 7

20 The following Graph A shows comparative tests results on a budesonide solution formulation comprising 0.22% w/w budesonide, 11% w/w ethanol, and the remainder 134a. Three lots were stored at 40°C/75% RH. One lot (identified as "Coated A") was contained in aluminum cans coated with an epoxy-phenolic coating (Cebal Printal Ltd. U.K.) and capped with a blind ferrule and a continuous gasket (made of DFDB 1085 elastomer, Union Carbide) so as to completely isolate the formulation from contact with
25 metal surfaces. Another lot ("Coated B") was contained in the same type of cans as Coated A, but equipped with a functional valve ferrule (3M Neotechnic Spraymiser™ solution valve) having a solution gasket to partially prevent contact of the formulation with the underside of the valve ferrule. A third lot ("Uncoated") was contained in uncoated aluminum cans equipped with the same valves as for Coated B.

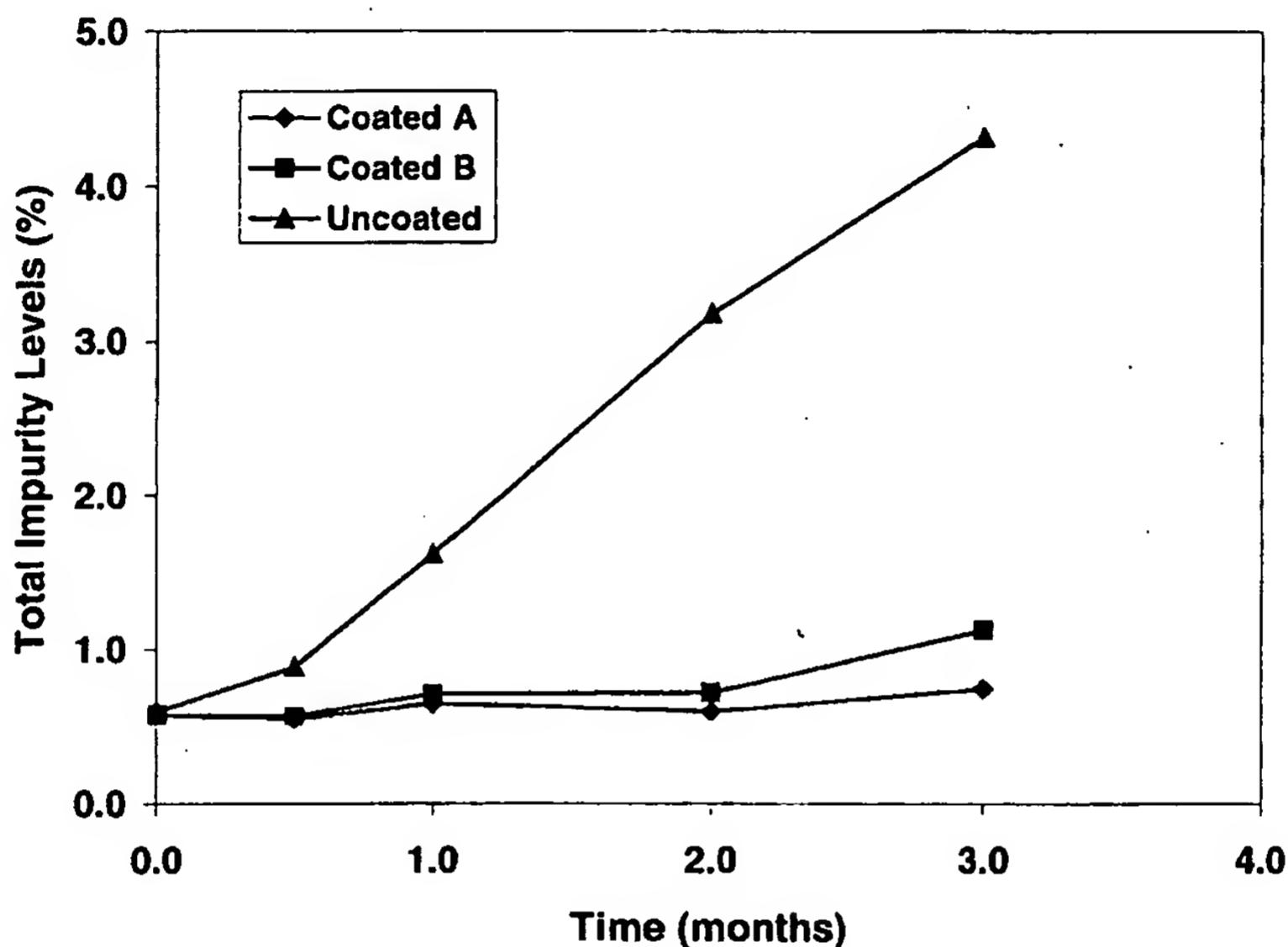
30 It can be seen from Graph A that the formulation in the Uncoated containers underwent much greater chemical degradation compared to the two coated containers, and that the Coated A lot underwent the least amount of degradation.

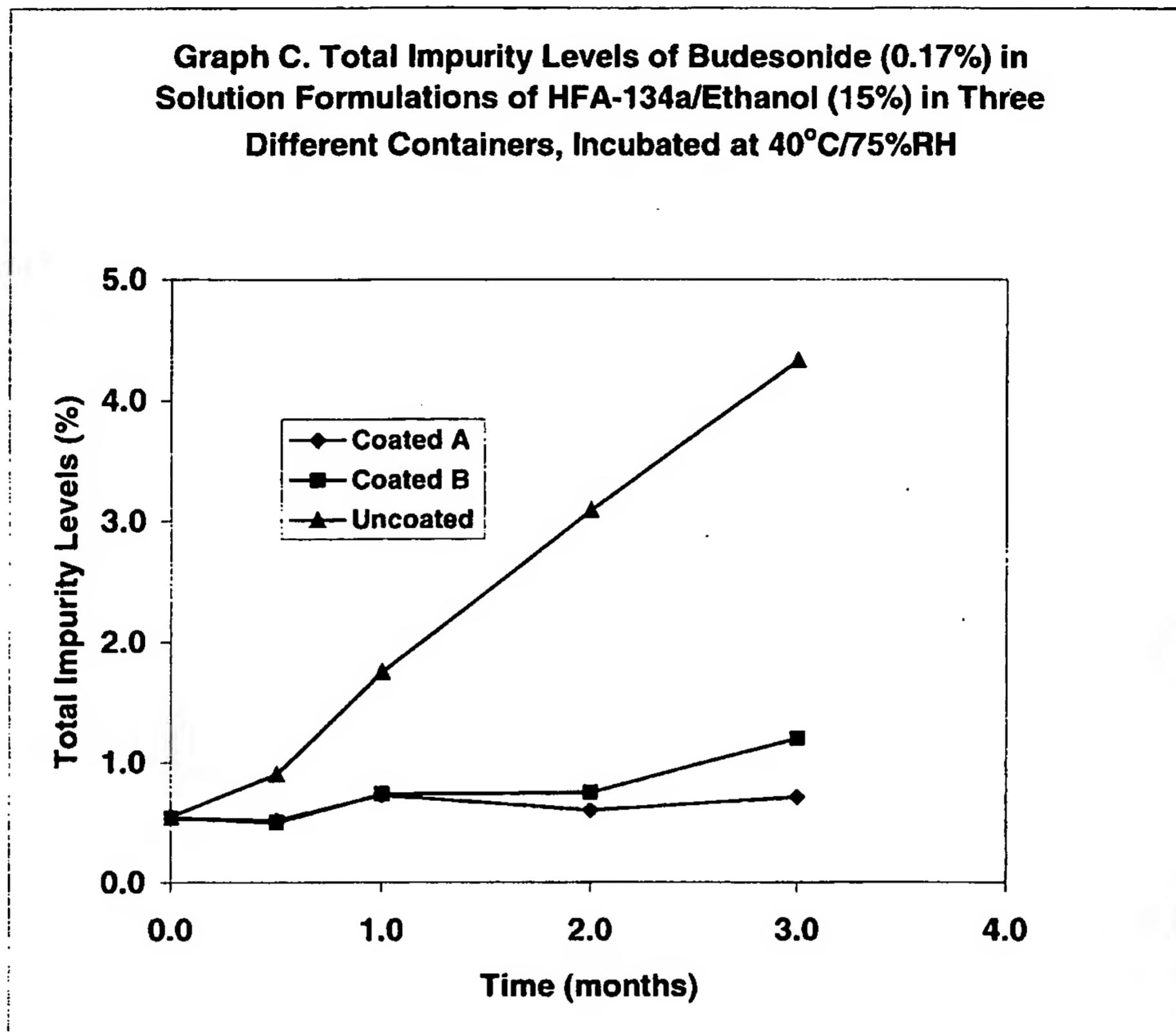


Examples 8-9

5 The same type of comparison (using the same kind of Coated A, Coated B, and
Uncoated containers as in example 7) was made for formulation Examples 8 and 9. In
Example 8 the formulation contained 0.22% w/w budesonide dissolved in a mixture of
134a and 13% w/w ethanol. The Example 9 formulation had 0.17% w/w budesonide
dissolved in 134a and 15% w/w ethanol. In both cases, shown in Graphs B and C,
10 respectively, the degradation rate was much greater in the Uncoated can, and degradation
was least for Coated A.

Graph B. Total Impurity Levels of Budesonide (0.22%) in Solution Formulations (HFA-134a/Ethanol 13%) in Three Different Containers, Incubated at 40°C/75%RH





Example 10

5 Test was also conducted on solution formulations of about 0.2% w/w budesonide, 13% anhydrous ethanol, and 134a to determine the effect of different coating types on chemical stability. Two types of canister coatings were compared: phenolic epoxy coating (Cebal) vs. an FEP coating (teflon-like polymer) from DuPont. Coated and uncoated valves similar to the design of Fig. 2 were compared: uncoated valves ("UV") 10 vs. coated valves ("CV") with Restek's SilcoSteel coated parts. The interior and exterior surfaces of the valve stem, metering tank, and bottle emptier were made of stainless steel and were coated with a submicron layer of fused silica. The results are summarized in Tables 6 and 7 below.

Table 6.

Container Closure System	Initial Testing*	1 month @40°C/75%RH
	Total Impurities %w/w	Total Impurities %w/w
UV/Epoxy lot A	1.09	0.98
CV/FEP lot B	1.17	1.36
UV/FEP lot C	1.36	2.31
CV/Epoxy lot D	1.17	0.95

Table 7.

Container Closure System	Initial Testing*	1 month @40°C/75%RH
	% Degradation Compared to Standard	% Degradation Compared to Standard
UV/Cebal lot A	0.03	0.17
CV/FEP lot B	0.11	0.55
FV/FEP lot C	0.30	1.50
UV/Epoxy lot D	0.11	0.14

5 *Note: the units were made and stored at ambient for about 2 months before "initial" testing. They were then placed in 40°C/75%RH chamber for stability. Thus, the initials are actually about 2 months @ ambient and the 1 month @ 40°C/75%RH is in addition to that period.

10 The results in Tables 6 and 7 indicate (1) that the coated valves (CV) improve chemical stability, and (2) that the epoxy coated canisters result in lower impurities than the FEP coating. These differing results based on the coating type are interesting because the FEP coating should provide equal protection to the formulation and, since the extractables profile of the FEP is believed to be much cleaner than the epoxy canister, the
15 FEP would have been expected to result in lower impurities.

 However, the data indicates the epoxy can is superior to the FEP in preventing degradation. Without wishing to be held to any particular mechanism to explain the difference in results between the two coating types, it is hypothesized that it is not that the phenolic epoxy provides a better barrier, but rather that some adsorption of degradation products is taking place. Essentially, the phenolic epoxy may be "soaking up" the degradates. A small amount of budesonide may also be adsorbed too, but the amount is apparently too small to affect the target content amount.
20

The description and examples set forth above are intended to illustrate the invention, but are not intended to be limiting. To the contrary, it will be understood that the invention is intended to include all variations and modifications falling within the scope of the appended claims. Also, although the invention is specifically applicable to C-5 17/21 20-ketosteroids in solution, it will be understood that particular steroid structures may have variations that modify the IUPAC carbon position numbering, thereby resulting in a position number other than 17 and/or 21 for OH groups that are nonetheless at effectively the same relative location. For example, if one were to remove the C-19 carbon atom, the numbering would change such that C-21 would be renamed C-20. Such 10 variations are intended to be within the scope of the invention and thus reference to "C-17" and "C-21" is intended herein to refer to the relative position based on the generic structures given above.

All percentages given herein are on a weight/weight (w/w) basis unless otherwise indicated. Also, all of the publications referred to herein are hereby incorporated by 15 reference.

CLAIMS

1. A medicinal aerosol steroid solution formulation product with enhanced chemical stability, comprising:
 - 5 an aerosol container equipped with a dispensing valve and containing a medicinal aerosol formulation having a 20-ketosteroid drug dissolved therein; said 20-ketosteroid having an OH group at the C-17 or C-21 position or both, with the proviso that said 20-ketosteroid is other than flunisolide; and
 - 10 wherein said container is provided with a non-metal interior surface so as to reduce chemical degradation of the 20-ketosteroid.
2. The product of claim 1, wherein said container is made of aluminum having an inert interior coating.
 - 15 3. The product of claim 2, wherein the interior coating is an epoxy-phenolic lacquer.
 4. The product of claim 1, wherein said dispensing valve is a metered dose valve.
- 20 5. The product of claim 1, wherein said medicinal aerosol formulation includes a hydrogen-containing propellant.
 6. The product of claim 5, wherein the hydrogen-containing propellant is a hydrofluorocarbon.
 - 25 7. The product of claim 6, wherein the hydrofluorocarbon propellant is selected from the group consisting of 1,1,1,2-tetrafluoroethane, 1,1,1,2,3,3,3-heptafluoropropane, and mixtures thereof.
 - 30 8. The product of claim 1, wherein said medicinal aerosol formulation includes ethanol.

9. The product of claim 1, wherein said 20-ketosteroid has an OH group at the C-17 position, but not at the C-21 position.
10. The product of claim 1, wherein said 20-ketosteroid has an OH group at the C-21 position, but not at the C-17 position.
5
11. The product of claim 1, wherein said 20-ketosteroid has an OH group at both the C-17 and C-21 positions.
- 10 12. The product of claim 1, wherein the 20-ketosteroid is a corticosteroid selected from the group consisting of budesonide, triamcinolone acetonide, desonide, fluocinolone acetonide, alclometasone, beclomethasone, beclomethasone 17-monopropionate, betamethasone, betamethasone 17-valerate, clocortolone, desoximetasone, dexamethasone, dexamethasone sodium phosphate, 15 dexamethasone 21-isonicotinate, diflorasone, flumethasone, methylprednisolone, paramethasone, prednisolone, triamcinolone, clobetasol, and fluorometholone.
13. The product of claim 1, wherein the 20-ketosteroid is budesonide.
- 20 14. The product of claim 1, wherein the 20-ketosteroid is triamcinolone acetonide.
15. The product of claim 1, wherein the 20-ketosteroid is dexamethasone.
16. The product of claim 1, wherein the 20-ketosteroid is betamethasone 17-valerate.
25
17. The product of any preceding claim wherein the container and/or the valve has a coating applied by vapor deposition.
18. The product of claim 17, wherein metal valve components have a coating applied by vapor deposition.
30
19. The product of claims 17 or 18, wherein the coating is a glass.

20. The product of claim 19, wherein the coating is applied by the Silcosteel process.
21. A method of reducing the chemical degradation of a medicinal 20-ketosteroid dissolved in a formulation contained in a metal container, said 20-ketosteroid being other than flunisolide and having an OH group at the C-17 position or C-21 position or both, comprising the step of providing a coating of inert material on the interior surface of the metal container so as to reduce reaction of the 20-ketosteroid with metal oxides from the container.
5
22. A process for making a chemically stable steroid solution aerosol product by filling into a container an aerosol formulation comprising a dissolved 20-ketosteroid other than flunisolide, said 20-ketosteroid having an OH group at the C-17 position or C-21 position or both, and said container having an inert non-metal interior surface so as to avoid chemical degradation of the 20-ketosteroid due to interaction with the container.
15
23. An aerosol valve having a non-metal coating applied to one or more metal surfaces by vapor deposition.
20
24. The valve of claim 23 wherein the coating is fused silica.
25. A medicinal aerosol product equipped with a metering valve and containing a medicinal aerosol formulation, said product comprising a layer of fused silica material applied on an internal surface of said product in contact with the formulation.
25
26. The product of claim 25, wherein said layer is on at least one component of the metering valve.
30
27. The product of claim 25, wherein said layer is sub-micron in thickness.

28. A method of making an improved metered dose medicinal aerosol product containing a medicinal aerosol formulation, comprising the step of coating an internal surface of said product, that is to be in contact with the formulation, with a layer of fused silica.

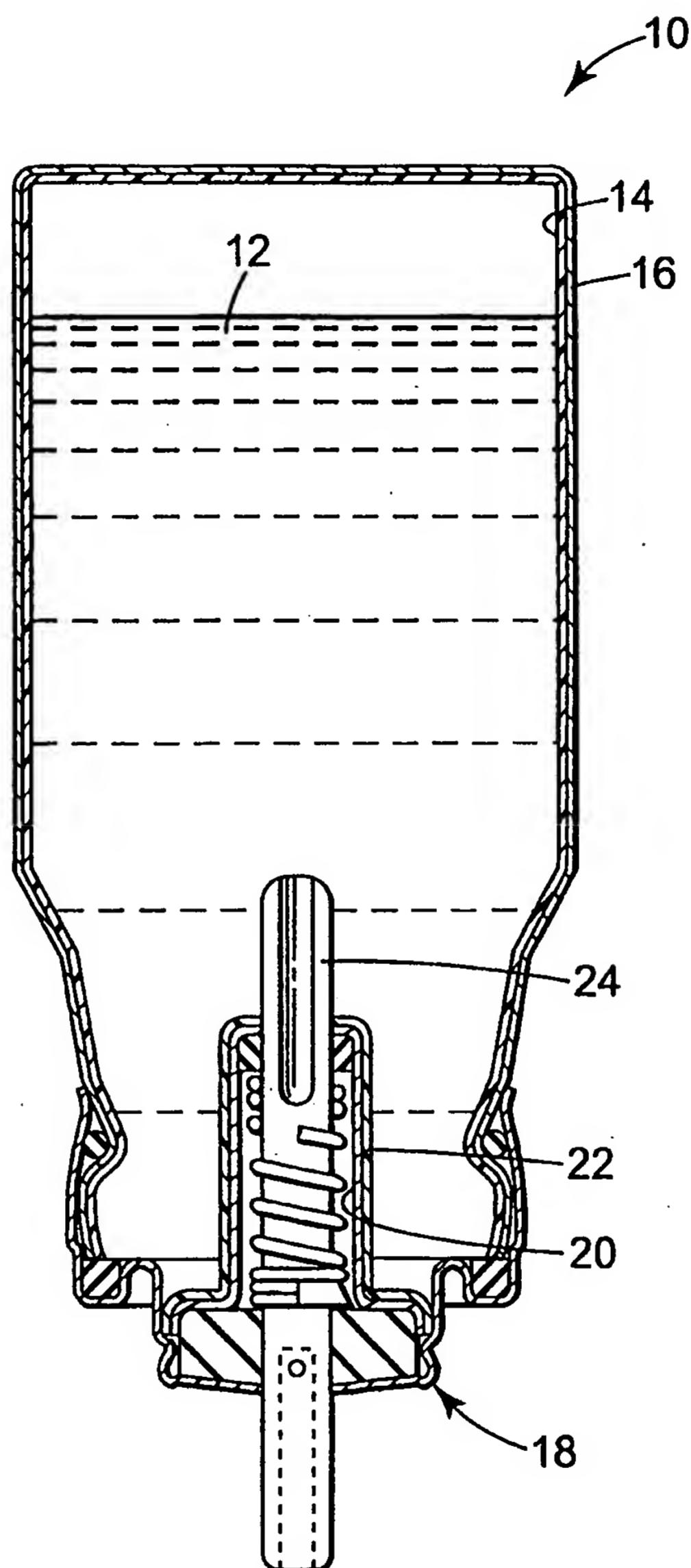


Fig. 1

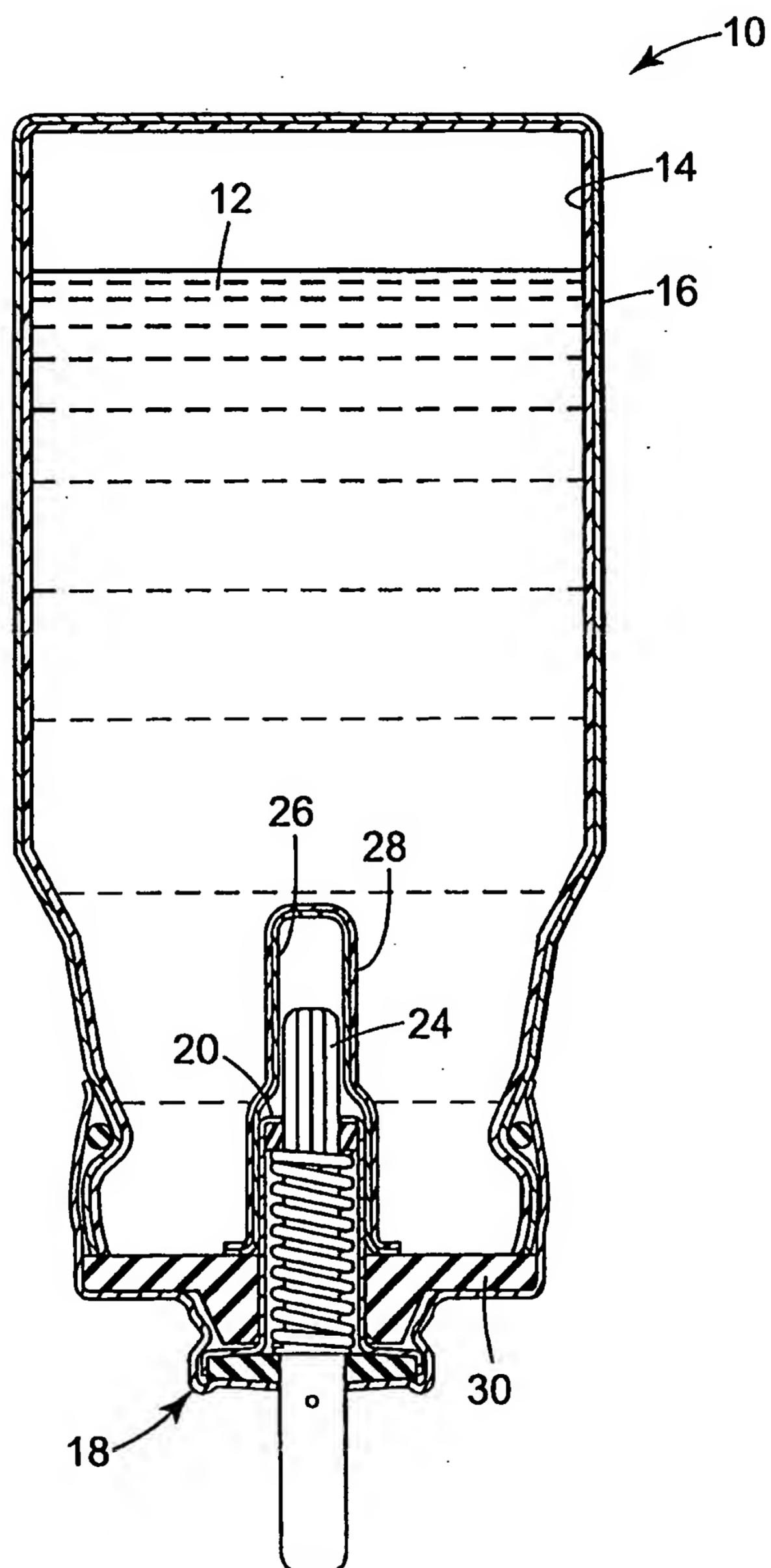


Fig. 2

INTERNATIONAL SEARCH REPORT

Internat' Application No

PCT/US 00/16462

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K9/12 B65D83/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 B65D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 776 433 A (TZOU TSI-ZONG ET AL) 7 July 1998 (1998-07-07) cited in the application column 2, line 35 - line 19 column 2, line 56 -column 3, line 2 column 4, line 1 - line 14 examples 19,20,29; tables 5,6,8 --- -/-/	1-8, 10, 21,22

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

° Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

10 October 2000

Date of mailing of the international search report

23/10/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Marttin, E

INTERNATIONAL SEARCH REPORT

Internal Application No
PCT/US 00/16462

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 98 24420 A (GOODMAN MICHAEL ;BIOGLAN IRELAND R & D LTD (IE); MCCARTHY PAUL (IE) 11 June 1998 (1998-06-11)</p> <p>page 2, line 21 -page 3, line 16 page 4, line 11 - line 13 page 5, line 19 -page 6, line 2 page 7, line 5 - line 8 page 8, line 7 - line 17 page 9, line 20 page 10, line 5 - line 11; figure 1 claims 1-3,11,17-20,22-32,34-40; examples 1-3</p> <p>---</p>	1,2,4-8, 11,12, 21,22
P,X	<p>WO 00 30608 A (BRAMBILLA GAETANO ;LEWIS DAVID (IT); VENTURA PAOLO (IT); GANDERTON) 2 June 2000 (2000-06-02)</p> <p>page 2, line 27 -page 3, line 6; claims 1-3,5-9 page 4, line 25 -page 5, line 28 page 8, line 19 - line 24; example 3; tables 4-9</p> <p>---</p>	1-8,10, 12-14, 19,21,22
P,X	<p>WO 99 47195 A (GLAXO GROUP LTD ;RIEBE MICHAEL THOMAS (US); SCHULZE MARK DOUGLAS () 23 September 1999 (1999-09-23)</p> <p>page 2, last paragraph -page 3, paragraph 1 page 3, last paragraph -page 4, paragraph 2 page 6, last paragraph page 7, paragraph 3 page 9, paragraph 2 page 10, paragraph 2; claims</p> <p>---</p>	23
X	US 5 743 251 A (SWEENEY WILLIAM R ET AL) 28 April 1998 (1998-04-28)	28
A	<p>column 1, line 7 - line 19 column 3, line 1 - line 6 column 3, line 11 - line 15 column 3, line 25 - line 40 column 11, line 12 - line 19 column 11, line 30 - line 40; claims 1,4,5,8-10</p> <p>---</p>	25-27
A	<p>"New SilcoCan canister with pressure/vacuum gauge" RESTEK ADVANTAGE CHROMATOGRAPHY NEWSLETTER, 'Online! 1996, XP002149698 Retrieved from the Internet: <URL:http://www.restekcorp.com/advntage/c96seven.htm> 'retrieved on 2000-10-09! the whole document</p> <p>-----</p>	1,17-20, 23-28

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/16462

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
US 5776433	A 07-07-1998	US 5980867 A		09-11-1999
		AU 680967 B		14-08-1997
		AU 1098695 A		10-07-1995
		CA 2178473 A		29-06-1995
		DE 69424181 D		31-05-2000
		EP 0735884 A		09-10-1996
		ES 2145254 T		01-07-2000
		JP 9506896 T		08-07-1997
		NO 962585 A		18-06-1996
		NZ 276637 A		27-07-1997
		WO 9517195 A		29-06-1995
-----	-----	-----	-----	-----
WO 9824420	A 11-06-1998	AU 5402898 A		29-06-1998
		EP 1011646 A		28-06-2000
		IE 970858 A		12-08-1998
		NO 992677 A		15-07-1999
		ZA 9710923 A		02-09-1998
-----	-----	-----	-----	-----
WO 0030608	A 02-06-2000	AU 1556300 A		13-06-2000
-----	-----	-----	-----	-----
WO 9947195	A 23-09-1999	AU 3145799 A		11-10-1999
-----	-----	-----	-----	-----
US 5743251	A 28-04-1998	AU 3205897 A		05-12-1997
		BR 9709252 A		10-08-1999
		CA 2255899 A		20-11-1997
		CZ 9803663 A		16-06-1999
		EP 0957959 A		24-11-1999
		NO 985265 A		14-01-1999
		PL 329869 A		12-04-1999
		WO 9742993 A		20-11-1997
-----	-----	-----	-----	-----